ABSTRACT

Described in the present invention are a sulfonyl derivative represented by the following formula (I):

$$O^1 - O^2 - T^1 - O^3 - SO_2 - O^A$$
 (I)

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[wherein Q¹ represents a saturated or unsaturated 5- or 6-membered cyclic hydrocarbon group, 5- or 6-membered heterocyclic group, dicyclic fused ring or tricyclic fused ring group which may have a substituent;

 Q^2 represents a single bond, an oxygen atom, a sulfur atom, a linear or branched C_{1-6} alkylene group or the like;

Q^A represents an arylalkenyl group which may have a substituent or a heteroarylalkenyl group which may have a substituent; and

T¹ represents a carbonyl group or the like] and a medicament comprising the same. The compound has strong FXa inhibitory action, provides prompt, sufficient and long-lasting anti-thrombus effects when orally administered, and has low side effects and is therefore useful as an excellent anticoagulant.